Effectiveness of www.SnelBeter.nl: activating occupational care on-line in employees with sickness absence due to back or neck pain.

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The purpose of this study is to investigate the effectiveness of a website. If the results indicate that the website is effective, the website may be implemented on a large scale. With the use of an effective website, companies will receive better...

Ethical review Approved WMO

Status Pending

Health condition type Muscle disorders **Study type** Interventional

Summary

ID

NL-OMON29814

Source

ToetsingOnline

Brief title

Effectiveness of www.SnelBeter.nl

Condition

· Muscle disorders

Synonym

back and neck disorders / pain, non specific back and neck complaints

Research involving

Human

Sponsors and support

Primary sponsor: Vrije Universiteit Medisch Centrum

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Source(s) of monetary or material Support: EMGO Instituut, VU medisch centrum: cofinandier voor de personele kosten tbv uitvoering en begeleiding van het onderzoek,KLM: cofinancier voor de directe kosten van de internet interventie en de uitvoering van interventies door arbo-professionals op basis van advies uit de internet interventie,NS: cofinancier voor de directe kosten van de internet interventie en de uitvoering van interventies door arbo-professionals op basis van advies uit de internet interventie,STECR Aladdin,WorkWell: cofinancier voor de kosten tbv de ontwikkeling van de website

Intervention

Keyword: activating occupational health care, effectiveness, internet, musculoskeletal disorders

Outcome measures

Primary outcome

There are three primary study parameters:

- 1) Employees understanding of their situation and autonomy
- 2) Performance of OPs
- 3) Time to refer by the OP

Secondary outcome

The secondary study parameter is time to full return to work.

Study description

Background summary

Many employees who are sicklisted do not really know how to handle the situation. Simple and specific education and individually based instructions through a website can possibly solve or reduce this problem. Besides, the website contains a questionnaire which offers relevant information for the occupational physician (OP). This can be a helpfull tool, because OPs unfortenately often neglet guidelines. Previous studies show that OPs do not refer al all or refer too late to specific interventions. This study will investigate the effectiveness of an interactive website. The hypothesis is that employees, by using this website, learn how to handle the situation. This will improve recovery and stimulate earlier return to work. Besides, the OPs will receive more information by using this website. Due to this extra information

OPs can refer quickly to an intervention.

Study objective

The purpose of this study is to investigate the effectiveness of a website.

If the results indicate that the website is effective, the website may be implemented on a large scale. With the use of an effective website, companies will receive better care from their occupational health service. This will result in a reduction of symptoms, disorders, sickness absence and costs.

Study design

Randomized Controlled Trial (RCT) with one intervention (n=64) and one control group (n=64). Randomisation will take place on the level of the occupational physician (OP). Furthermore, the prerandomisation design will be used. This means that participants are not aware of the randomisation procedure, but will only be informed about the general purpose of the study.

Intervention

All participants will receive usual care in accordance with the guidelines for OPs.

Besides, participants in the intervention group will also use the website. This website is an extra tool to understand their situation and illustrates what they can do themselves to solve the problem. The website generates simple but specific information about the situation for each employee. Moreover, the website gives individually based instructions.

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Study burden and risks

At two moments, participants are asked to fill in a questionnaire. We estimated that one questionnaire will take 20 minutes to fulfill. Participants in the intervention group will also need some time to read the information on the website en perform the instructions which are given. It is difficult to estimate the amount of time needed for the website.

The are no risks involved in this study.

All participants receive usual care in accordance with the guidelines for OPs. Participants in the intervention group will also follow instructions on the website, which can not lead to serious adverse events. All exercises are based on the nature and history of the complaints.

Moreover, specific or serious complaints will be checked by the OP before

someone can participate in the study.

Contacts

Public

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

Non specific back or neck complaints and sickness absence due to these complaints Understanding of the Dutch language Internet access also at home

Exclusion criteria

Serious complaints (red flags, e.g. high fever), which need immediately medical care from a

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Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Single blinded (masking used)

Control: Active

Primary purpose: Treatment

Recruitment

NL

Recruitment status: Pending

Start date (anticipated): 01-09-2006

Enrollment: 128

Type: Anticipated

Ethics review

Approved WMO

Application type: First submission

Review commission: METC Amsterdam UMC

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

CCMO NL11705.029.06